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INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 6 : A61K 38/04, 38/08, 38/09, 38/24, 47/08, 47/16, 47/18, 47/20, 47/02		A1	(11) International Publication Number: WO 98/00157 (43) International Publication Date: 8 January 1998 (08.01.98)
(21) International Application Number: PCT/US97/10816 (22) International Filing Date: 1 July 1997 (01.07.97) (30) Priority Data: 60/021,199 3 July 1996 (03.07.96) US (60) Parent Application or Grant (63) Related by Continuation US 60/021,199 (CIP) Filed on 3 July 1996 (03.07.96) (71) Applicant (for all designated States except US): ALZA CORPORATION [US/US]; 950 Page Mill Road, P.O. Box 10950, Palo Alto, CA 94303-0802 (US). (71) Applicant (for US only): ECKENHOFF, Bonnie, J. (legal representative of the deceased inventor) [US/US]; 1080 Autumn Lane, Los Altos, CA 94024 (US). (72) Inventor: ECKENHOFF, James, B. (deceased). (72) Inventors; and (75) Inventors/Applicants (for US only): STEVENSON, Cynthia, L. [US/US]; 100 West El Camino Real #48, Mountain View,		CA 94040 (US). TAO, Sally, A. [US/US]; 1151 Miller Avenue, San Jose, CA 95129 (US). PRESTRELSKI, Steven, J. [US/US]; 1971 West Middlefield Road #5, Mountain View, CA 94043 (US). WRIGHT, Jeremy, C. [US/US]; 631 Cuesta Drive, Los Altos, CA 94024 (US). LEONARD, Joe [US/US]; 11236 La Jolla Court, Cupertino, CA 95014 (US). (74) Agents: DHUEY, John, A. et al.; Alza Corporation, 950 Page Mill Road, P.O. Box 10950, Palo Alto, CA 94303-0802 (US). (81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ARIPO patent (GH, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i> <i>Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>	
(54) Title: AQUEOUS FORMULATIONS OF PEPTIDES			
(57) Abstract <p>This invention relates to stable liquid aqueous formulations of peptide compounds at high concentrations. These stable formulations comprise at least about 10 % peptide in water. They may be stored at elevated temperatures for long periods of time and are especially useful in implantable delivery devices for long term delivery of drug.</p>			

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AQUEOUS FORMULATIONS OF PEPTIDES

5

FIELD OF THE INVENTION

This invention relates to stable aqueous formulations of peptide
10 compounds at high concentrations.

BACKGROUND OF THE INVENTION

References:

The following references are referred to by numbers in brackets ([]) at
15 the relevant portion of the specification.

1. Zoladex (goserelin acetate implant), Physician's Desk Reference, 50th Edition, pages 2858-2861 (1996).
2. U.S. Patent No. 3,914,412, issued October 21, 1975.
3. U.S. Patent No. 4,547,370, issued October 15, 1985.
- 20 4. U.S. Patent No. 4,661,472, issued April 28, 1987.
- 5 U.S. Patent No. 4,689,396, issued August 25, 1987.
- 6 U.S. Patent No. 4,851,385, issued July 25, 1989.
7. U.S. Patent No. 5,198,533, issued March 30, 1993.
8. U.S. Patent No. 5,480,868, issued January 2, 1996.
- 25 9. WO92/20711, published 26 November 1992.
10. WO95/00168, published 5 January 1995.
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12. "Stability of Gonadorelin and Triptorelin in Aqueous Solution", V.J. Helm, B.W. Muller, *Pharmaceutical Research*, 7/12, pages 1253-1256
30 (1990).
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15. "Parenteral Peptide Formulations: Chemical and Physical Properties of Native Luteinizing Hormone-Releasing Hormone (LHRH) and Hydrophobic Analogues in Aqueous Solution", M.F. Powell, L.M. Sanders, A. Rogerson, V. Si, *Pharmaceutical Research*, 8/10, pages 1258-1263 (1991).
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17. "Percutaneous Absorption Enhancement of Leuprolide", M.Y. Fu Lu, D. Lee, G.S. Rao, *Pharmaceutical Research*, 9/12, pages 1575-1576 (1992).
18. Lutrepulse (gonadorelin acetate for IV injection), Physician's Desk Reference, 50th Edition, pages 980-982 (1996).
19. Factrel (gonadorelin HCl for subcutaneous or IV injection), Physician's Desk Reference, 50th Edition, pages 2877-2878 (1996).
20. Lupron (leuprolide acetate for subcutaneous injection), Physician's Desk Reference, 50th Edition, pages 2555-2556 (1996).
21. Lupron depot (leuprolide acetate for depot suspension), Physician's Desk Reference, 50th Edition, pages 2556-2562 (1996).
22. "Pharmaceutical Manipulation of Leuprorelin Acetate to Improve Clinical Performance", H. Toguchi, *J. of Intl. Medical Research*, 18, pages 35-41 (1990).
23. "Long-Term Stability of Aqueous Solutions of Luteinizing Hormone-Releasing Hormone Assessed by an In-Vitro Bioassay and Liquid Chromatography", Y.F. Shi, R. J. Sherins, D. Brightwell, J.F. Gallelli, D.

C. Chatterji, *J. of Pharmaceutical Sciences*, 73/6, pages 819-821 (1984).

24. "Peptide Liquid Crystals: Inverse Correlation of Kinetic Formation and Thermodynamic Stability in Aqueous Solution", M.F. Powell, J. Fleitman, L.M. Sanders, V.C. Si, *Pharmaceutical Research*, 11/9, pages 1352-1354 (1994).

25. "Solution Behavior of Leuprolide Acetate, an LHRH Agonist, as Determined by Circular Dichroism Spectroscopy", M.E. Powers, A. Adejei, M.Y. Fu Lu, M.C. Manning, *Intl. J. of Pharmaceutics*, 108, pages 49-55 (1994).

26. "Preparation of Three-Month Depot Injectable Microspheres of Leuprorelin Acetate Using Biodegradable Polymers", *Pharmaceutical Research*, 11/8, pages 1143-1147 (1994).

The disclosure of each of the above publications, patents or patent applications is hereby incorporated by reference in its entirety to the same extent as if the language of each individual publication, patent and patent application were specifically and individually incorporated by reference.

Luteinizing hormone-releasing hormone (LHRH), also known as gonadotropin releasing hormone (GnRH), is a decapeptide with the structure:

pGlu-His-Trp-Ser-Tyr-Gly-Leu-Arg-Pro-Gly-NH₂.

It is secreted by the hypothalamus and binds to receptors on the pituitary gland, releasing luteinizing hormone (LH) and follicle stimulating hormone (FSH). LH and FSH stimulate the gonads to synthesize steroid hormones. Numerous analogs of LHRH are known, including peptides related to LHRH which act as agonists and those which act as antagonists. [1-15] LHRH analogs are known to be useful for treating hormone-dependent diseases such as prostate cancer, benign prostatic hyperplasia, endometriosis, hysteromyoma, metrorrhagia, precocious puberty, or mammary cancer and as contraceptives. [8] Sustained release administration is preferred for both agonist LHRH-related compounds, which reduce the number of available receptors after repeated administration so that the production of steroid hormones is suppressed, and antagonist LHRH-related compounds, which

must be continually administered for persistent inhibition of endogenous LHRH. [8]

The sustained parenteral delivery of drugs, especially peptide drugs, provides many advantages. The use of implantable devices for sustained
5 delivery of a wide variety of drugs or other beneficial agents is well known in the art. Typical devices are described, for example, in U.S. Patents Nos. 5,034,229; 5,057,318; and 5,110,596. The disclosure of each of these patents is incorporated herein by reference.

In general, oral bioavailability of peptides, including LHRH-related
10 compounds, is low. [16-17]

Currently marketed aqueous formulations of LHRH, its analogs and related compounds which are used for parenteral injection generally contain relatively low concentrations of LHRH-related compounds (0.05 to 5 mg/ml) and may also contain excipients such as mannitol or lactose. [18-20] Such
15 formulations of LHRH-related compounds must either be stored refrigerated or may be stored at room temperature for short periods of time.

Available depot formulations of LHRH-related compounds administered for sustained release over a period of 1-3 months include a formulation comprised of 15% LHRH-related compound dispersed in a matrix
20 of D,L-lactic and glycolic acids copolymer presented as a cylinder to be injected subcutaneously [1] and a formulation comprised of microparticles comprising a core of LHRH-related compound and gelatin surrounded by a shell of D,L-lactic and glycolic acids copolymer. These microparticles are suspended in a diluent for injection either subcutaneously or intramuscularly.
25 [21, 26] These products must be stored at room temperature or lower. Aqueous formulations of LHRH-related compounds are known to exhibit both chemical and physical instability, as well as degradation after irradiation. [12-16, 22-25]

Formulations which have been shown to be stable (t_{90} about five years)
30 have been very low concentration (25 μ g/ml) aqueous, buffered (10 mM buffer, ionic strength of 0.15) solutions stored at temperatures no higher than room temperature (25°C). [15]

There is a need for stable, high concentration aqueous formulations of peptides.

SUMMARY OF THE INVENTION

5 The present invention provides stable aqueous formulations which are solutions of peptides compounds in water at concentrations of at least about 10%. These stable high concentration formulations may be stored at elevated temperatures (e.g., 37°C) for long periods of time and are especially useful in implantable delivery devices for long term delivery (e.g., 1-12
10 months or longer) of drug. The aqueous formulations may optionally include buffer, excipients, ethanol (EtOH), a surfactant or a preservative.

In one aspect, the invention provides stable aqueous formulations of peptide compounds, said formulations comprising at least about 10% (w/w) peptide compound and water.

15 In another aspect, the invention provides methods for preparing a stable aqueous formulation of an peptide compound, said methods comprising dissolving at least about 10% (w/w) peptide compound in water.

In yet a further aspect, the invention provides methods for treating a subject suffering from a condition which may be alleviated by administration
20 of a peptide compound, said methods comprising administering to said subject an effective amount of a stable aqueous formulation comprising at least about 10% (w/w) peptide compound and water.

BRIEF DESCRIPTION OF THE DRAWINGS

25 Figure 1 illustrates the stability of 40% leuprolide acetate solution in water after two months at 80° C as measured by reverse phase HPLC (RP-HPLC).

Figure 2 shows the same sample as Figure 1 injected by size exclusion chromatography (SEC). This figure shows that there is very little aggregation, and what aggregation there is is comprised of dimer and trimer products, with
30 no higher order aggregation.

Figure 3 presents the Arrhenius plot showing the loss of leuprolide from 40% solutions of leuprolide acetate in water.

Figure 4 illustrates the chemical and physical stability of a 40% leuprolide acetate solution in water after about three months at 80°C.

5 Figure 5 illustrates the loss of leuprolide acetate fit to pseudo first order kinetics from a 40% solution in water over a period of three to six months at 37°C, 50°C, 65°C and 80°C.

Figure 6 illustrates the chemical and physical stability of a 40% leuprolide acetate solution in water after nine months at 37°C.

10 Figure 7 illustrates the stability of a 30% goserelin solution in acetate buffer and mannitol after 14 days at 80°C.

Figure 8 illustrates that both gelled and non-gelled aqueous formulations of leuprolide (370 mg/ml) remained stable over a period of 6 months at 37°C.

15

DETAILED DESCRIPTION OF THE INVENTION

The present invention is drawn to the unexpected discovery that dissolving high concentrations (i.e., at least about 10%) of peptide compounds in water results in stable aqueous formulations. Previously
20 known aqueous formulations of peptide compounds, which are dilute buffered aqueous solutions containing excipients such as EDTA or ascorbic acid which must be stored at low temperatures (4-25°C), form degradation products using degradation pathways such as acid/base catalyzed hydrolysis, deamidation, racemization and oxidation. In contrast, the presently claimed
25 formulations stabilize peptide compounds at high concentrations at elevated temperatures (e.g., 37°C to 80°C), thus making possible the delivery of peptides in implantable delivery devices that would not otherwise be feasible.

Standard peptide and protein formulations consist of dilute aqueous solutions. Two critical aspects of peptide formulation include solubilization
30 and stabilization of the drug molecule. Peptide stability is usually achieved by varying one or more of the following: pH, buffer type, ionic strength, excipients

(EDTA, ascorbic acid, etc.). In contrast, in the present invention, highly concentrated peptides formulated in water provide stable solutions.

The invention consists of using high concentrations of peptide in aqueous solution to stabilize the peptide formulations against both chemical
5 and physical degradation.

A. Definitions:

As used herein, the following terms have the following meanings:

The term "chemical stability" means that an acceptable percentage of
10 degradation products produced by chemical pathways such as oxidation or hydrolysis is formed. In particular, a formulation is considered chemically stable if no more than about 20% breakdown products are formed after two months at 37°C.

The term "physical stability" means that an acceptable percentage of
15 aggregates (e.g., dimers, trimers and larger forms) is formed. In particular, a formulation is considered physically stable if no more than about 15% aggregates are formed after two months at 37°C.

The term "stable formulation" means that at least about 65%
20 chemically and physically stable peptide compound remains after two months at 37°C (or equivalent conditions at an elevated temperature). Particularly preferred formulations are those which retain at least about 80% chemically and physically stable peptide under these conditions. Especially preferred stable formulations are those which do not exhibit degradation after sterilizing irradiation (e.g., gamma, beta or electron beam).

25 The terms "peptide" and/or "peptide compound" mean polymers of up to about 50 amino acid residues bound together by amide (CONH) linkages. Analogs, derivatives, agonists, antagonists and pharmaceutically acceptable salts of any of these are included in these terms. The terms also include peptides and/or peptide compounds which have D-amino acids, modified,
30 derivatized or non-naturally occurring amino acids in their D- or L- configuration and/or peptomimetic units as part of their structure.

The term "LHRH-related compound" means luteinizing hormone releasing hormone (LHRH) and its analogs and pharmaceutically acceptable salts. Octa-, nona- and decapeptide LHRH agonists and antagonists are included in the term LHRH-related compounds, as is native LHRH.

5 Particularly preferred LHRH-related compounds include LHRH, leuprolide, goserelin, nafarelin, and other known active agonists and antagonists. [1-21]

The term "high concentration" means at least about 10% (w/w) and up to the maximum solubility of the particular LHRH-related compound.

The term "excipient" means a more or less inert substance in a
10 formulation which is added as a diluent or vehicle or to give form or consistency. Excipients are distinguished from solvents such as EtOH, which are used to dissolve drugs in formulations, from non-ionic surfactants such as Tween 20, which are used to solubilize drugs in formulations, and from
15 preservatives such as benzyl alcohols or methyl or propyl parabens, which are used to prevent or inhibit microbial growth.

The term "buffering capacity" means the capacity of a solution due to the presence of a mixture of an acid/base pair in the solution to reduce any changes in pH that would otherwise occur in the solution when acid or alkali is added to it.

20 The term "polar aprotic solvent" means a polar solvent which does not contain acidic hydrogen and does not act as a hydrogen bond donor. Examples of polar aprotic solvents are dimethylsulfoxide (DMSO), dimethylformamide (DMF), hexamethylphosphorotriamide (HMPT), and n-methyl pyrrolidone.

25

B. Preparation of Formulations:

The present invention is drawn to highly concentrated liquid aqueous formulations of peptide compounds which are stable for prolonged periods of time at elevated temperatures. Standard dilute aqueous peptide and protein
30 formulations require manipulation of buffer type, ionic strength, pH and excipients (e.g., EDTA and ascorbic acid) to achieve stability. In contrast, the claimed formulations achieve stabilization of peptide compounds by the use

of high concentrations (at least about 10%, w/w) of compound dissolved in water.

Examples of peptides and peptide compounds which may be formulated using the present invention include those peptides which have biological activity or which may be used to treat a disease or other pathological condition. They include, but are not limited to adrenocorticotrophic hormone, angiotensin I and II, atrial natriuretic peptide, bombesin, bradykinin, calcitonin, cerebellin, dynorphin A, alpha and beta endorphin, endothelin, enkephalin, epidermal growth factor, fertirelin, follicular gonadotropin releasing peptide, galanin, glucagon, gonadorelin, gonadotropin, goserelin, growth hormone releasing peptide, histrelin, insulin, leuprolide, LHRH, motilin, nafarelin, neurotensin, oxytocin, somatostatin, substance P, tumor necrosis factor, triptorelin, and vasopressin. Analogs, derivatives, antagonists, agonists and pharmaceutically acceptable salts of the above may also be used.

Depending on the particular peptide compound to be formulated, ionic strength and pH may be factors worthy of consideration. For example, we have found that preferred aqueous formulations of leuprolide acetate have low ionic strength and pH between about 4 and about 6.

The peptide compounds useful in the formulations and methods of the present invention can be used in the form of a salt, preferably a pharmaceutically acceptable salt. Useful salts are known to those of skill in the art and include salts with inorganic acids, organic acids, inorganic bases or organic bases. Preferred salts are acetate salts.

Peptide compounds which are hydrophilic and readily soluble in water are preferred for use in the present invention. One of skill in the art can easily determine which compounds will be useful on the basis of their aqueous solubility, i.e., the compound must be soluble in water to at least about 10% (w/w). Preferably, this is also a pharmaceutically effective amount.

Particularly preferred peptide compounds are LHRH-related compounds, including leuprolide and leuprolide acetate.

The proportion of peptide may vary depending on the compound, the condition to be treated, the solubility of the compound, the expected dose and the duration of administration. (See, for example, The Pharmacological Basis of Therapeutics, Gilman et al., 7th ed. (1985) and Pharmaceutical Sciences, Remington, 18th ed. (1990), the disclosures of which are incorporated herein
5 by reference.) The concentration of peptide compound may range from at least about 10% (w/w) to the maximum solubility of the compound. A preferred range is from about 20 to about 60% (w/w). The currently more preferred range is from about 30 to about 50% (w/w) and a most preferred
10 range is about 35 to about 45% (w/w).

Generally, the stable formulations of the present invention may be prepared by simply dissolving a therapeutically effective amount of the desired peptide compound in water, although pH adjustments may be made.

It is known to those of skill in the art that buffers, excipients, solvents
15 such as EtOH, solubilizers such as non-ionic surfactants, and preservatives may beneficially be added to pharmaceutical peptide formulations. (See, for example, Pharmaceutical Sciences, Remington, 18th ed. (1990).) Such agents may optionally be added to the claimed formulations.

20 C. Methodology:

We have found that stable aqueous formulations of peptide compounds may be prepared by dissolving a high concentration (at least about 10%) of the peptide compound to be formulated in water.

We have tested these peptide compound formulations, specifically
25 formulations of the LHRH-related compound leuprolide, for stability by subjecting them to accelerated aging at elevated temperature and measuring the chemical and physical stability of the formulations. Results of these studies (shown, for example, in Table III and Figures 1, 2 and 6) demonstrate that these formulations were stable at conditions that approximate or exceed
30 storage for one year at 37°C.

We have also tested peptide compound formulations prepared as described herein for stability after 2.5 megarad gamma irradiation. Results,

shown in Table IV, show that these formulations remained chemically and physically stable after such irradiation. Formulations subjected to electron beam irradiation were also found to be stable.

As shown in Table I, we have tested a wide variety of peptide
5 formulations, specifically leuprolide, goserelin, LHRH, angiotensin I,
bradykinin, calcitonin, insulin, trypsinogen and vasopressin, for stability by
dissolving (or attempting to dissolve) them in water, then subjecting them to
accelerated aging at elevated temperatures. The stability of the formulations
was measured. Results are presented in Table I as half-life at 37°C
10 assuming an $E_a = 22.2$ kcal/mole. A wide range of the peptides tested were
soluble in water and remained stable under the test conditions. The solubility
of a particular peptide in water and the stability of the resulting solution are
easily determined using routine procedures known to those of ordinary skill in
the art.

Table I: Stability of Peptides Formulated in Water

FORMULATION	HALF-LIFE* (Temperature)
40% Leuprolide	9.7 years (37°C)
40% Goserelin	19.3 months (80°C)
20% LHRH	2.5 years (65°C)
20% Angiotensin I	insoluble gel (65°C)
20% Bradykinin	8.5 months (65°C)
40% Calcitonin	insoluble (80°C)
20% Calcitonin	9.6 months (80°C)
5% Calcitonin	23.5 months (50°C)
20% Insulin	insoluble gel (65°C)
40% Trypsinogen	insoluble gel (65°C/80°C)
20% Trypsinogen	insoluble gel (65°C)
40% Vasopressin	degraded (80°C)
20% Vasopressin	14.3 days (65°C)
*Half-life at 37°C assuming $E_a = 22.2$ kcal/mole.	

Formulations of 40% leuprolide in water stored for six months at 37°C showed linear degradation as measured by overall loss of peptide from the solution. Analysis of these data gave an activation energy (E_a) of 22.2 kcal/mole and a t_{90} of 13.8 months, showing stability of these formulations at elevated temperatures.

We have also unexpectedly found that certain peptide formulations of the present invention are bacteriostatic (i.e., inhibit bacterial growth), bactericidal (i.e., cause the death of bacteria), and sporicidal (i.e., kill spores). In particular, leuprolide formulations of 50-400 mg/ml exhibited bacteriostatic, bactericidal and sporicidal activity. The stability of the samples was unaffected by spiking with bacteria, indicating that the enzymes released from

the killed and lysed bacteria did not adversely affect the stability of the product. This demonstrates that these formulations were not conducive to enzymatic activity.

Some peptides, for example calcitonin and leuprolide, are known to be physically unstable, exhibiting aggregation, gelation and fibrillation when formulated in aqueous solution. For example, leuprolide can be induced to gel by increasing peptide concentration, introduction of salts or gentle agitation. Improving physical stability can allow for easier parenteral administration, including administration using implantable drug delivery systems.

It has unexpectedly been found that adding polar aprotic solvents such as DMSO to aqueous formulations of certain peptides, such as leuprolide, goserelin and calcitonin, prevents gelation of the formulation. This is apparently because non-aqueous polar aprotic solvents cause peptides to form a random coil/alpha helix conformation that does not refold into a beta sheet structure and, therefore, does not gel. Thus, these solvents have an anti-gellant effect.

Additionally, studies of gelled and non-gelled aqueous formulations of leuprolide (370 mg/ml) stored at 37°C for 6 weeks showed a similar chemical stability profile as assayed by RP-HPLC. Results are shown in Figure 8. Similarly, the stability of liquid and gelled (by agitation) aqueous leuprolide formulations (370 mg/ml) was studied in vitro at 37°C and in vivo in rats, respectively. Results are presented in Table II, and show that the both gelled and liquid formulations remained stable over a period of 18 weeks.

Table II: Stability Studies of Liquid and Gelled Aqueous L-leuprolide Formulations

STUDY	TIME (weeks)	LIQUID (% remaining)	GELLED (% remaining)
Long Term Stab	6	98.00	
Long Term Stab	12	91.50	
Long Term Stab	18	93.50	
Rat	4		94.80
Rat	6		93.50
Rat	12		92.30
Rat	18		92.60

A major aspect of the invention is that aqueous solutions containing
5 high concentrations of peptide compounds are stable at high temperatures for
long periods of time. Thus, these formulations are advantageous in that they
may be shipped and/or stored for long periods of time at or above room
temperature. They are also suitable for use in implantable delivery devices.

DISCLOSURE OF EXAMPLES OF THE INVENTION

The following methods were used to perform the studies in the Examples that follow.

5 1. Preparing leuprolide acetate solutions

Leuprolide acetate (obtained, for example, from Mallinckrodt, St. Louis, Missouri) was weighed, added to a weighed amount of vehicle (sterile distilled water, ethanol/water or water with non-ionic surfactant) at the appropriate concentration (w/w), then gently stirred to dissolve.

10 Unless otherwise noted, leuprolide free base content was calculated from certificate of analysis potency values to be 37% free base. This was 40% leuprolide acetate, except as noted.

 2. Preparation of reservoirs

15 The reservoirs of implantable drug delivery devices (as disclosed in U.S. Patent Application Serial No. 08/595,761, incorporated herein by reference) were filled with the appropriate leuprolide acetate solution. The filled devices then underwent stability testing. The formulation was filled into titanium or polymer reservoirs with a polymer plug blocking each end. The
20 filled reservoir was then sealed in a polyfoil bag and placed in a stability testing oven.

It should be noted that the formulations in the reservoirs of these devices are completely isolated from the outside environment.

25 3. Reverse Phase-HPLC (RP-HPLC)

All stability samples were analyzed for leuprolide concentration and % peak area using a gradient elution reversed-phase HPLC assay with a refrigerated autosampler (4°C) to minimize sample degradation. The chromatographic conditions used are listed below.

RP-HPLC Chromatographic Conditions

Description	Parameter																		
Column	HaiSil C18, 4.6 X 250mm, S/N 5103051																		
Flow Rate	0.8 mL min ⁻¹																		
Injection Volume	20 µL																		
Detection	210 nm																		
Leuprolide Retention Time	Between 25-30 minutes																		
Mobile Phase	A = 100 mM Sodium Phosphate, pH 3.0 B = 90% Acetonitrile/Water																		
Gradient	<table><tr><td>Minutes</td><td>0</td><td>5</td><td>25</td><td>40</td><td>41</td><td>46</td><td>46.1</td><td>50</td></tr><tr><td>%B</td><td>15</td><td>26.5</td><td>26.5</td><td>65</td><td>85</td><td>85</td><td>15</td><td>15</td></tr></table>	Minutes	0	5	25	40	41	46	46.1	50	%B	15	26.5	26.5	65	85	85	15	15
Minutes	0	5	25	40	41	46	46.1	50											
%B	15	26.5	26.5	65	85	85	15	15											

Leuprolide standards (in water) at 4 to 6 different concentration levels, typically between 0.1 - 1.2 mg/mL, were run along with the stability samples.

- 5 The stability samples were bracketed by the standard sets, with no more than 40 samples in between the standard sets. All peaks between the void volume and 45 minutes of the run were integrated. The integrated peak areas for the leuprolide standards were plotted as a function of the concentration. The leuprolide concentrations for the stability samples were then calculated using
- 10 linear regression. The % peak areas for the leuprolide peak, the sum of all the peaks eluting before leuprolide (labeled "others"), and the sum of all the peaks eluting after leuprolide (labeled "aggregates") were also recorded and plotted as a function of the sample timepoints.

15 4. Size Exclusion Chromatography (SEC)

Selected stability samples were analyzed for % peak area and molecular weights using an isocratic solution SEC assay with a refrigerated autosampler (4°C). The chromatographic conditions used are listed below.

SEC Chromatographic Conditions

Description	Parameter
Column	Pharmacia Peptide, HR 10/30, 10 X 300 mm
Flow Rate	0.5 mL min ⁻¹
Injection Volume	20 µL
Detection	210 nm
Leuprolide Retention Time	Approximately 25 minutes
Mobile Phase	100 mM Ammonium Phosphate, pH 2.0, 200 mM Sodium Chloride, 30% Acetonitrile

The void volume and total volume for the size exclusion column was needed for the calculation of the molecular weights. The BioRad high molecular weight standard and 0.1% acetone were used to determine the void volume and total volume respectively. The retention times for the first peak in the BioRad standard and the acetone peak were recorded and converted to volume units using the equations below. Since these values are constant for a particular SEC column and HPLC system, the void and total volumes were redetermined whenever changes to the SEC column or HPLC system were made. A standard run was then made followed by the stability samples. The standard mixture contained approximately 0.2 mg/mL of the following peptides: Bursin (MW=449), WLFR peptide (MW=619), Angiotensin (MW=1181), GRF (MW=5108), and Cytochrome C (MW=12394). These standards were chosen because they bracketed leuprolide molecular weight and all had basic pI (9.8 - 11.0), similar to leuprolide.

The % peak areas were recorded for all the peaks. The molecular weights for the species separated were calculated using the equations below.

$$V_s = \text{flow rate (mL/min)} \times \text{sample peak retention time (min)}$$

$$V_o = \text{flow rate (mL/min)} \times \text{void volume peak retention time (min)}$$

$$V_t = \text{flow rate (mL/min)} \times \text{total volume peak retention time (min)}$$

$$K_d = \frac{V_s - V_o}{V_t - V_o}$$

The following examples are offered to illustrate this invention and are not meant to be construed in any way as limiting the scope of this invention.

EXAMPLE 1

Accelerated Stability Studies of Leuprolide Acetate Formulations.

5 Formulations of 40% (w/w) leuprolide acetate (equivalent to about 37% leuprolide free base) in either sterile distilled water, ethanol/water (70/30) or water with 10% Tween 20 were prepared as described above and used to fill the reservoirs of implantable drug delivery devices, also as described above.
10 Some reservoirs were made of polymer materials, while some were titanium.

 The filled devices were subjected to accelerated aging by storing them at elevated temperatures (80-88°C) for seven days in an incubator (Precision Scientific or Thelco). This is equivalent to about 1.5 years at 37°C or about four years at room temperature (25°C), assuming an activation energy (E_a) or
15 22.2 kcal/mole.

 The samples were analyzed using RP-HPLC and SEC as described above to determine the chemical and physical stability of the aged formulations.

 Results, presented in Table III, demonstrate that these aqueous
20 formulations were able to maintain the stability of the LHRH-related compound leuprolide. In each case, at least 65% leuprolide was retained. However, a large amount of the formulation with EtOH evaporated from the reservoir during the study, indicating that long term storage at elevated temperatures of formulations with high concentrations of a volatile solvent like
25 EtOH may be problematic. The formulation which contained the non-ionic surfactant 10% Tween 20 was found to be no more stable than water solutions without this solubilizer.

Table III**Stability of 40% (w/w) Leuprolide Acetate Aqueous Formulations After 7 Days at Elevated Temperatures**

Temperature (°C)	Reservoir Material	Formulation	% Leuprolide at Day 7
88	Polymer	40% in Water	68
88	Titanium	40% in Water	71
88	Polymer	40% in Water	66*
88	Polymer	40% in EtOH/H ₂ O (70/30)	85**
88	Polymer	40% in 10% Tween 20	65
80	Polymer	40% in Water	83
80	Polymer	40% in Water	80
80	Polymer	40% in Water	78
80	Polymer	40% in Water	79
80	Polymer	40% in Water	83
80	Polymer	40% in Water	77
80	Polymer	40% in Water	79
80	Polymer	40% in Water	74
80	Polymer	40% in Water	88

* 10% evaporated

** 60% evaporated

EXAMPLE 2**Stability Studies of Irradiated Leuprolide Acetate Formulations**

Formulations of 40% (w/w) as received leuprolide acetate (equivalent to 37% leuprolide free base) in water were prepared as described above and
5 used to fill the reservoirs of drug delivery devices, also as described above. Some reservoirs were made of polymer materials, while some were titanium.

The filled devices were subjected to 2.5 megarad gamma irradiation. Samples were shipped to Sterigenics (Tustin, California) and gamma irradiated (Cobalt 60) in batch mode. Samples were then subjected to
10 accelerated aging as in Example 1. Samples labeled "cold" were shipped and irradiated on dry ice. Samples were taken at day 0 and day 7, and analyzed using RP-HPLC and SEC as described above to determine the chemical and physical stability of the irradiated formulations.

Results, presented in Table IV, demonstrate that these leuprolide
15 acetate formulations were stable after irradiation. In every case, at least 65% leuprolide was retained, with low levels of aggregate formation.

Table IV
Stability of 40% (w/w) Leuprolide Acetate Aqueous Formulations After 2.5 Megarad Gamma Irradiation

Reservoir Material	Formulation	Irradiation	% Leuprolide at Day 7 (RP-HPLC)	SEC			
				Day 0		Day 7	
				% monomer	% dimer/trimer	% monomer	% dimer/trimer
Polymer	40 % in Water	Yes	75	90.4	1.2	80.9	3.9
Polymer	40 % in Water	No	75	99.8	0.2	82.4	3.1
Polymer	40 % in Water	Cold	79	89.4	0.2	80.3	3.1
Titanium	40 % in Water	Yes	83	98.5	1.1	84.9	2
Titanium	40 % in Water	No	N.D.	99.6	0	96.6	0
Titanium	40 % in Water	Yes	81	98.8	0.9	94.7	2.4
Titanium	40 % in Water	No	82	99.9	0	95	1.9
Titanium	40 % in Water	Yes	73	99.1	0.9	88.3	3
Titanium	40 % in Water	Yes	79	99	0.8	94.3	3.4
Titanium	40 % in Water	Yes	74	98.6	0.5	90.9	3.6

EXAMPLE 3

Long Term Accelerated Stability Studies of Leuprolide Acetate in Water

Solutions of 40 % leuprolide acetate (w/w) in water were prepared, loaded into reservoirs, stored for two months at 80°C and analyzed as described above. Results, shown in Figures 1 (RP-HPLC) and 2 (SEC) show that 81.1% leuprolide was recovered, with only 14.6% chemical degradation and 5.1% physical aggregation after the two month period.

Leuprolide acetate solutions (40% (w/w) in water) were prepared, loaded, stored and analyzed as set forth above. Figure 4 is a plot of leuprolide, and its chemical and physical degradation products recovered over a three month time period. The sum of these three elements is also presented as mass balance. The results show that we can account for all the peptide material as either intact leuprolide or a degradation species, indicating that stability studies are not missing an unknown degradation process or product.

Leuprolide acetate solutions (40% (w/w) in water) were prepared, loaded, stored at 37°C, 50°C, 65°C or 80°C and analyzed using RP-HPLC as described above. Figure 5 shows the loss of leuprolide from these solutions over a three to six month period, and indicates that leuprolide degradation fits pseudo first order kinetics. Furthermore, as discussed below, Figure 3 indicates that leuprolide in water degradation fits linear Arrhenius kinetics. Therefore, accelerated stability studies are a valid technique for assessing the stability of leuprolide and extrapolating back to 37°C.

Solutions of 40% leuprolide acetate (w/w) in water were prepared, loaded into reservoirs, stored at 37°C, 50°C, 65°C or 80°C and analyzed using RP-HPLC as described above. Results were calculated as described in Physical Pharmacy: Physical Chemical Principles in the Pharmaceutical Sciences, 3rd ed., Martin et al., Chapter 14 (1983) and showed the E_a of these solutions to be 22.2 kcal/mole with a t_{90} of 13.8 months.

The data are shown below and an Arrhenius plot of the data is presented in Figure 3.

5

Water		
°C	Kobs (months ⁻¹)	t _{1/2} (months)
37	7.24×10^{-3}	95.7
50	3.21×10^{-2}	21.6
65	0.111	6.3
80	0.655	1.1

 $E_a = 22.2 \text{ kcal/mole}$

EXAMPLE 4

Long Term Stability Studies of Leuprolide Acetate in Water

The chemical stability of 40% leuprolide acetate solutions prepared and analyzed as described above is presented in Figure 6. After nine months at 37°C more than 85% (88.3%) leuprolide was present, with less than 10% (8.4%) chemical degradation products (shown as "early" in the figure, based on the RP-HPLC profile) and less than 5% (3.5%) physical aggregates (shown as "late" in the figure, based on RP-HPLC data, but in good agreement with SEC data).

EXAMPLE 5

Accelerated Stability Studies of Goserelin

Formulations of 30% goserelin (w/w) in acetate buffer (pH 5.0, 0.0282M) with 3% mannitol were stored in glass ampules for 14 days at 80°C and analyzed for purity as described above.

Results in Figure 7 show that after 9 days about 65% goserelin remained.

EXAMPLE 6**Stability Studies of Goserelin Formulations**

Formulations of 40-45% (w/w) goserelin in either acetate buffer, with 3% mannitol or acetate buffer with salt (0.9% NaCl) were prepared as described above and placed in polymeric containers.

The containers were stored at 37°C for one month in an incubator.

The samples were analyzed using RP-HPLC to determine the chemical stability of the aged formulations.

Results, presented below, demonstrate that these aqueous formulations were able to maintain the stability of the LHRH-related compound goserelin. In each case, at least 98% goserelin was retained.

DRUG	VEHICLE	% PURITY	% CONCENTRATION
Goserelin	Acetate Buffer/Mannitol	98.1	54.2
Goserelin	Acetate Buffer/Salt	98.0	50.1

EXAMPLE 7**Stability Studies of Nafarelin Formulations**

Formulations of 15% (w/w) nafarelin in acetate buffer with 3% mannitol were prepared as described above and placed in polymeric containers.

The containers were stored at 37°C for one month in an incubator.

The samples were analyzed using RP-HPLC to determine the chemical stability of the aged formulations.

Results, presented below, demonstrate that these aqueous formulations were able to maintain the stability of the LHRH-related compound nafarelin, as at least 98% nafarelin was retained.

DRUG	VEHICLE	% PURITY	% CONCENTRATION
Nafarelin	Acetate Buffer/Mannitol	98.8	18.3

Modification of the above-described modes of carrying out various embodiments of this invention will be apparent to those of skill in the art

following the teachings of this invention as set forth herein. The examples described above are not limiting, but are merely exemplary of this invention, the scope of which is defined by the following claims.

What is claimed is:

1. A stable aqueous formulation of a peptide related compound comprising:
 - a) at least about 10% (w/w) of at least one peptide compound;and
 - b) water.
2. The formulation of Claim 1 which comprises at least about 30% (w/w) peptide compound.
- 15 3. The formulation of Claim 1 wherein said peptide compound is an LHRH-related compound.
4. The formulation of Claim 3 wherein said peptide compound is selected from the group consisting of leuprolide, LHRH, nafarelin and goserelin.
5. The formulation of Claim 1 which is stable after irradiation.
6. The formulation of Claim 1 which is stable at 80°C for at least 2 months.
7. The formulation of Claim 1 which is stable at 37°C for at least 3 months.
- 30 8. The formulation of Claim 1 which is stable at 37° C for at least one year.
9. The formulation of Claim 1 which is adapted for use in an implantable drug delivery device.

10. The formulation of Claim 1 which further comprises at least one selected from the group consisting of a buffer, an excipient, a solvent, a solubilizer and a preservative.

11. The formulation of Claim 1 which consists essentially of about 30% to about 50% (w/w) of the LHRH-related compound leuprolide acetate in sterile distilled water.

12. The formulation of Claim 1 which forms a gel.

13. The formulation of Claim 1 further comprising at least one non-aqueous polar aprotic solvent.

14. The formulation of Claim 13 wherein said non-aqueous polar aprotic solvent is DMSO or DMF.

20

15. A method for preparing the stable aqueous formulation of Claim 1 comprising dissolving at least about 10% (w/w) of at least one peptide compound in water.

16. The method of Claim 15 wherein at least about 30% (w/w) peptide compound is dissolved.

17. The method of Claim 15 wherein said peptide compound is an LHRH-related compound.

30

18. The method of Claim 17 wherein said peptide compound is selected from the group consisting of leuprolide, LHRH, nafarelin and goserelin.

19. The method of Claim 15 further comprising the step of adding at least one selected from the group consisting of a buffer, an excipient, a solvent, a solubilizer and a preservative.

20. The method of Claim 15 wherein about 30% to about 50% (w/w) of the LHRH-related compound leuprolide acetate is dissolved in sterile distilled water.

21. The method of Claim 15 further comprising the step of adding at least one non-aqueous polar aprotic solvent.

15

22. The method of Claim 21 wherein said non-aqueous polar aprotic solvent is DMSO or DMF.

23. A method for treating a subject suffering from a condition which may be alleviated by administration of an peptide compound comprising administering to said subject an effective amount of the formulation of Claim 1.

24. The method of Claim 23 wherein said administration is parenteral administration.

25. The method of Claim 23 wherein said administration is long-term continuous administration.

26. The method of Claim 25 wherein said administration is accomplished by use of an implantable drug delivery device.

27. The method of Claim 23 wherein said condition is prostatic cancer and said peptide compound is leuprolide.

28. The method of Claim 23 wherein at least about 80 micrograms of leuprolide is administered daily.

29. The method of Claim 28 wherein said daily administration continues for a period selected from the group consisting of 3 months, 6 months and 12 months.

30. The method of Claim 29 wherein said daily administration for said period is continuous administration accomplished using an implantable drug delivery system.

15

31. The method of Claim 23 wherein said condition is prostatic cancer and said peptide compound is an LHRH antagonist.

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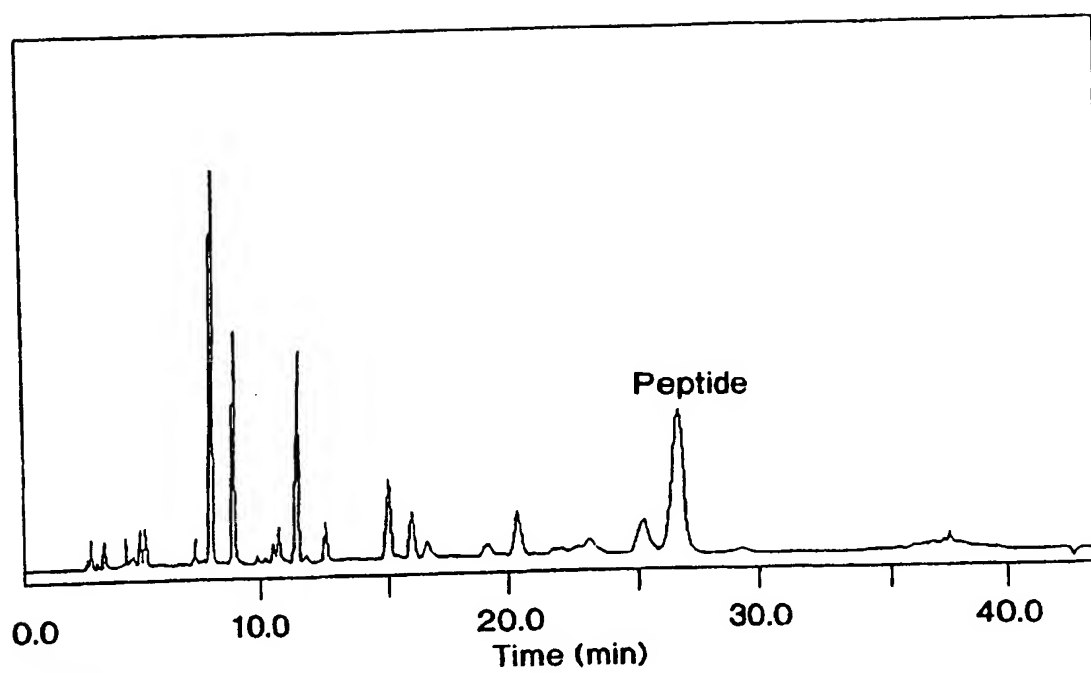


FIG. 1

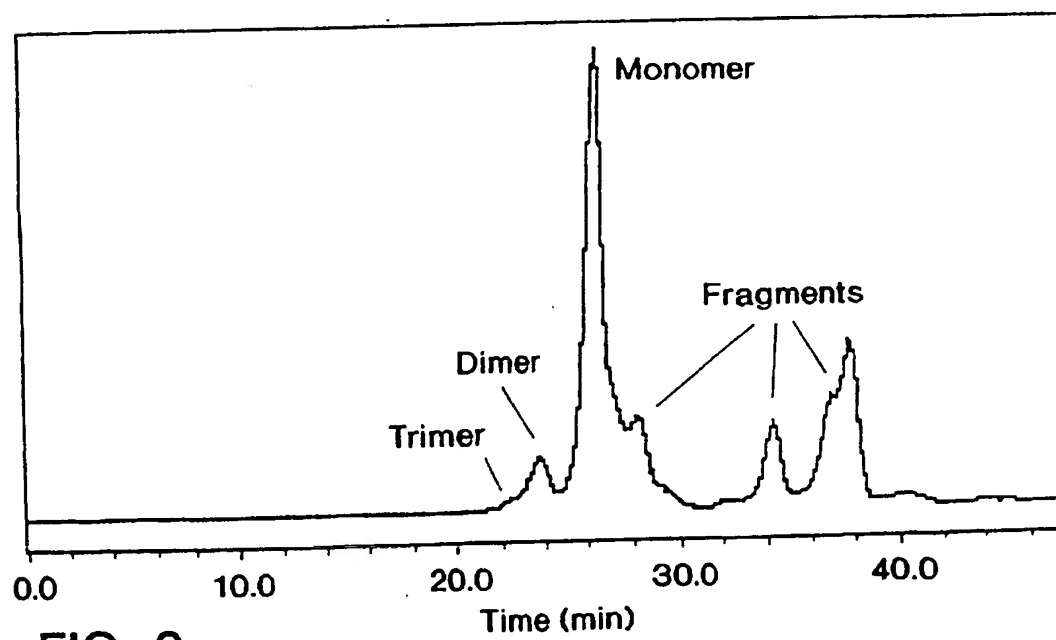


FIG. 2

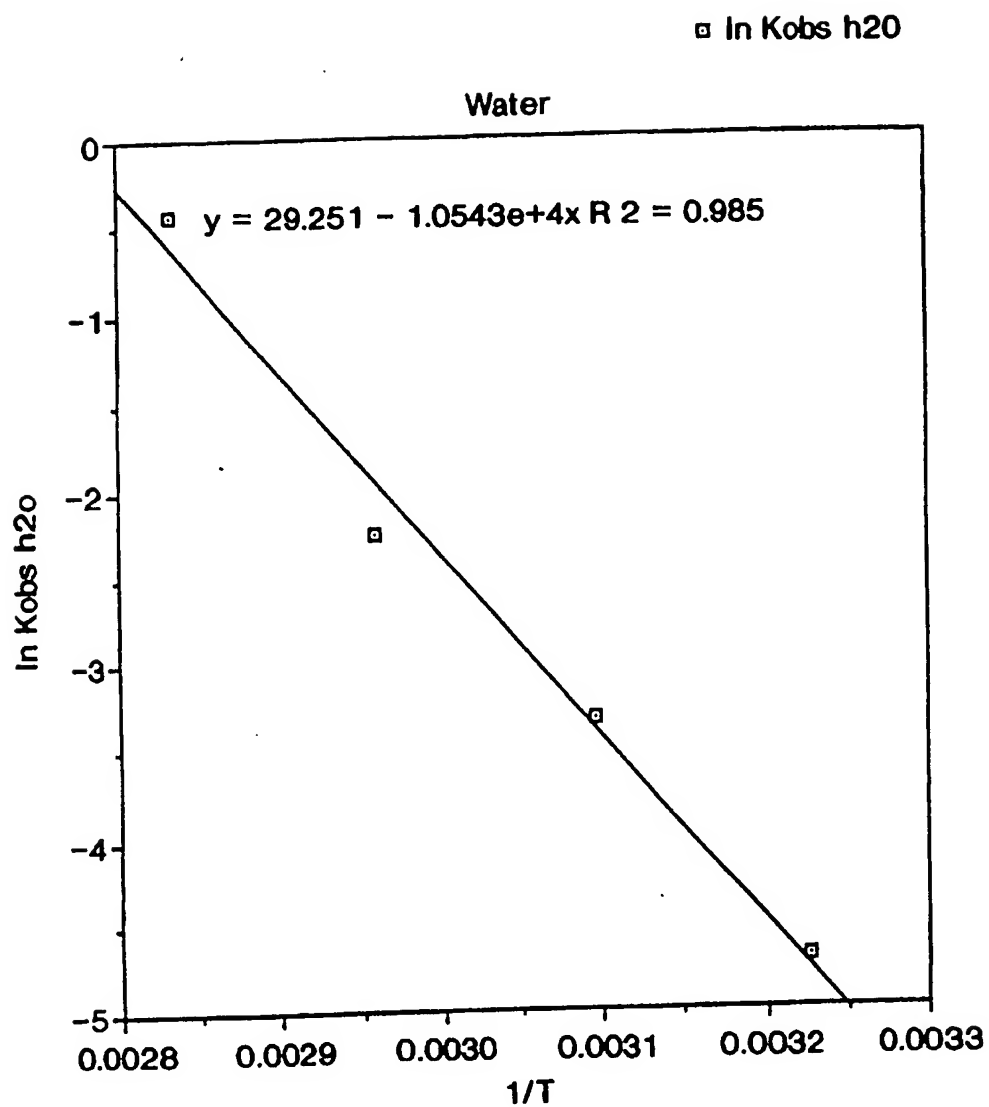


FIG. 3

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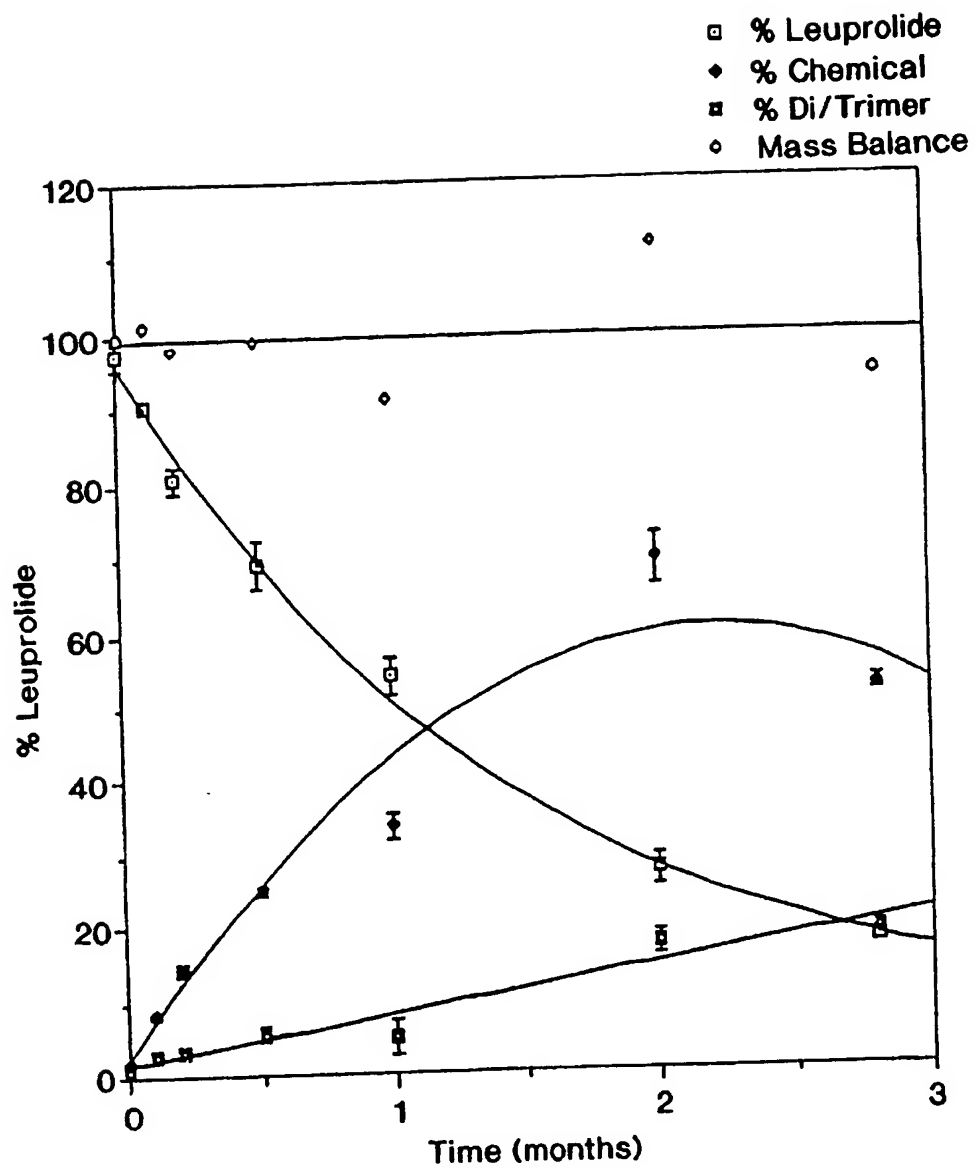


FIG. 4

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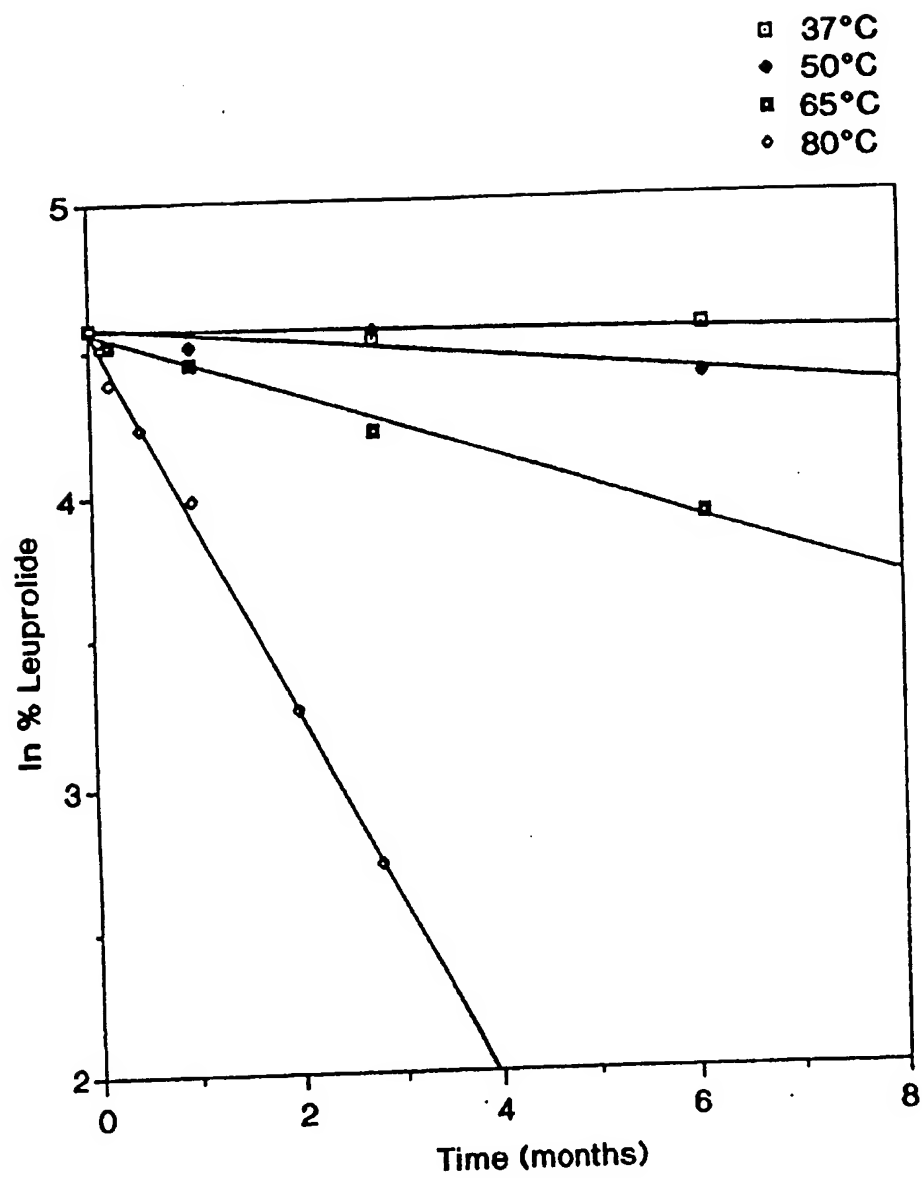


FIG. 5

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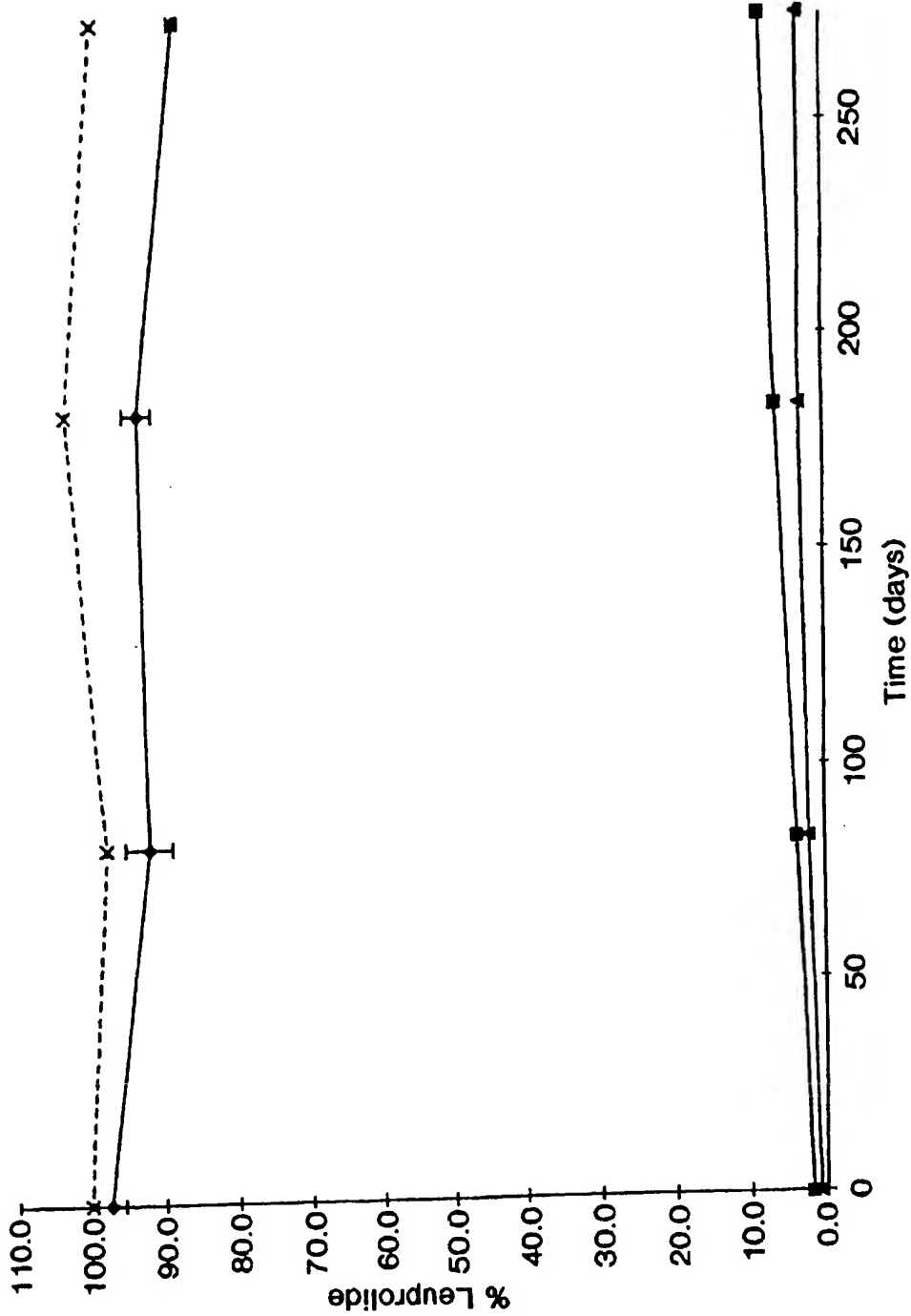


FIG. 6

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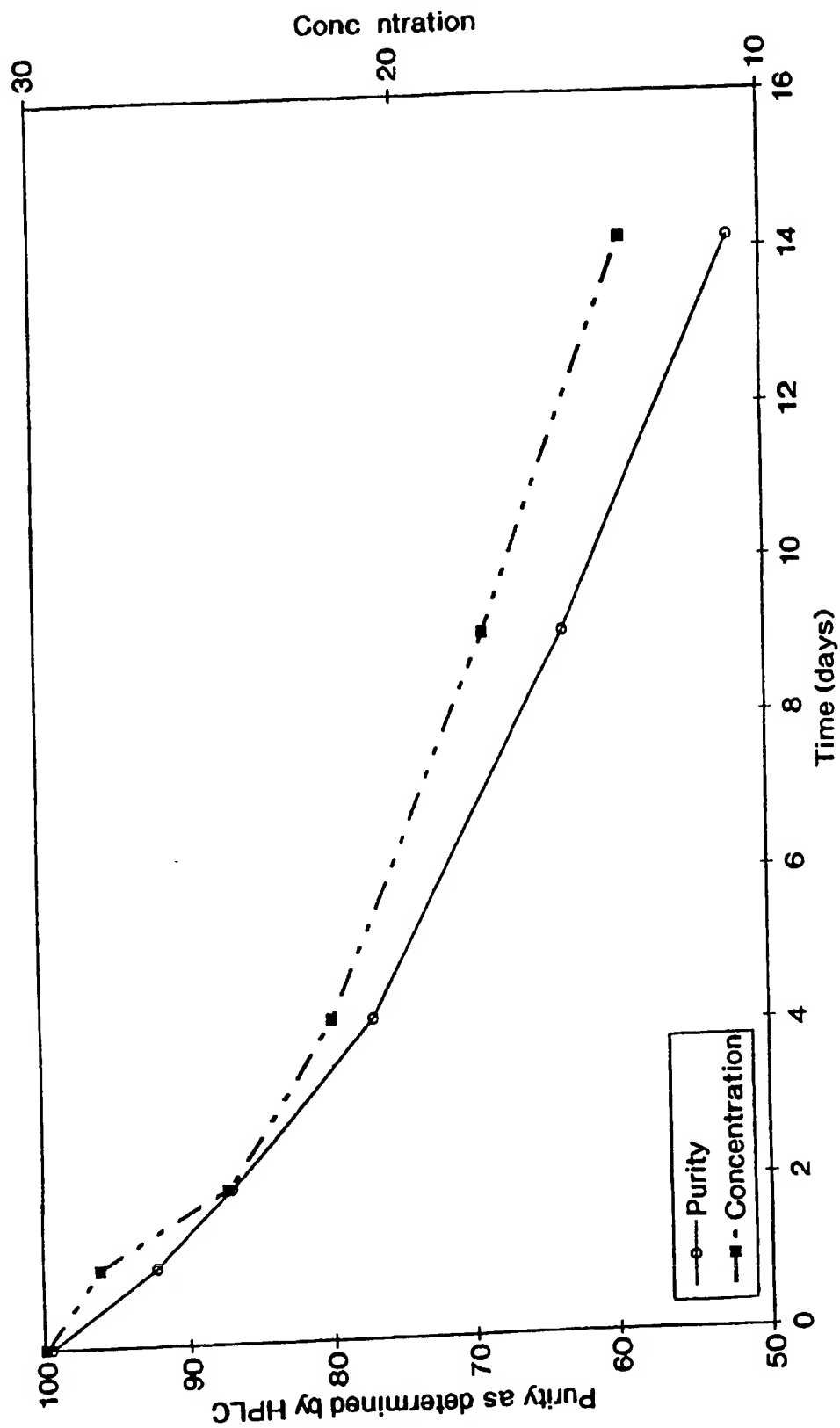


FIG. 7

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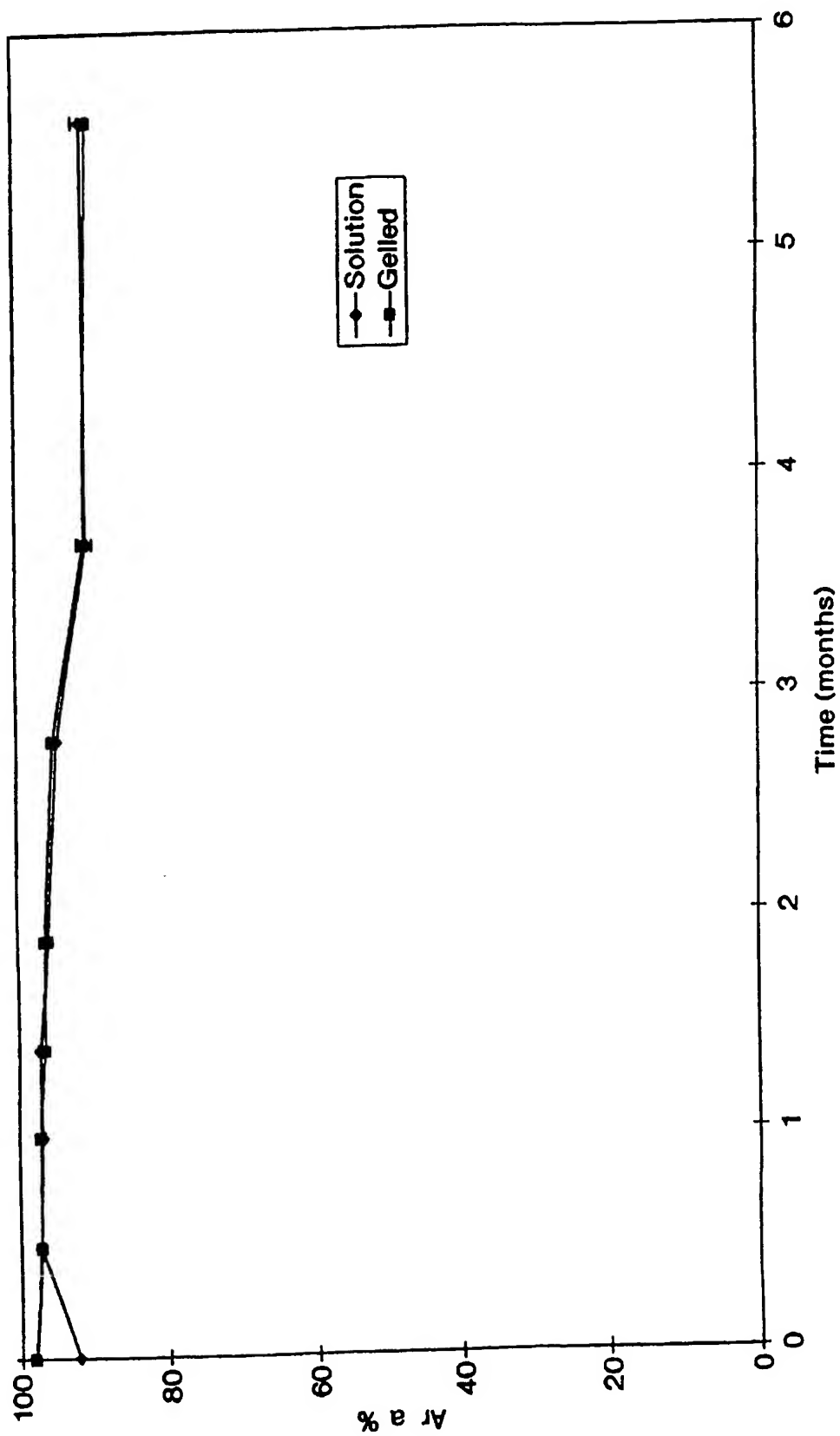


FIG. 8

INTERNATIONAL SEARCH REPORT

Intern. Application No
PCT/US 97/10816

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 A61K38/04 A61K38/08 A61K38/09 A61K38/24 A61K47/08
A61K47/16 A61K47/18 A61K47/20 A61K47/02

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 96 07398 A (SOD CONSEILS RECH APPLIC ;CHERIF CHEIKH ROLAND (FR)) 14 March 1996	1-10, 15-19, 23-26
Y	see the whole document	13,14, 21,22
X	EP 0 111 841 A (SYNTEX INC) 27 June 1984	1,3-8, 10,15, 17-19, 23-25
	see the whole document	
Y	WO 94 19020 A (GENENTECH INC ;CLELAND JEFFREY L (US); JONES ANDREW J S (US)) 1 September 1994 see the whole document	13,14, 21,22
A	WO 95 01183 A (FERRING AB) 12 January 1995	
	-/--	

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents :

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
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- *P* document published prior to the international filing date but later than the priority date claimed

T later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

X document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

Y document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

A document member of the same patent family

Date of the actual completion of the international search

26 November 1997

Date of mailing of the international search report

11. 12. 97

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INTERNATIONAL SEARCH REPORT

Intern. Application No
PCT/US 97/10816

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 5 482 931 A (HARRIS ALAN ET AL) 9 January 1996	
A	US 5 498 598 A (HARRIS ALAN) 12 March 1996	

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US 97/10816

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
see FURTHER INFORMATION sheet PCT/ISA/210
2. ☐ Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

Claims Nos.: 23-31

because they relate to subject matter not required to be searched by this Authority, namely:

Rule 39.1(iv) PCT - Method for treatment of the human or animal body by therapy

Remark : Although claims 24-31 are directed to a method of treatment of the human/animal body , the search has been carried out and based on the alleged effects of the compound/composition.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 97/10816

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INTERNATIONAL SEARCH REPORT

Information on patent family members

Intern. Appl. Application No

PCT/US 97/10816

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
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